

# Patent omphalomesenteric duct: sectioning the unexpected

Isabel Rosário Periquito,<sup>1</sup> Tânia Marques,<sup>2</sup> Sofia Lima,<sup>3</sup> Marta Ferreira<sup>2</sup>

<sup>1</sup>Department of Pediatrics, Centro Hospitalar de Setúbal, Setúbal, Portugal

<sup>2</sup>Department of Neonatology, Hospital Prof. Doutor Fernando Fonseca, Amadora, Portugal

<sup>3</sup>Department of Pediatric Surgery, Hospital Prof. Doutor Fernando Fonseca, Amadora, Portugal

## Correspondence to

Dr Isabel Rosário Periquito, isabelperiquito@gmail.com

Accepted 14 October 2014

## DESCRIPTION

We report the case of a male newborn, the second triplet of non-consanguineous healthy parents, born from a dichorionic triamniotic pregnancy with adequate prenatal care and normal ultrasounds. Spontaneous labour occurred at 32 weeks of gestation and a caesarean section was performed. The Apgar score was 5/6/8, birth weight was 1395 g and no malformations were noted. After sectioning the umbilical cord for venous catheter placement, an abnormal structure with a lumen discharging watery liquid was noted (figure 1). The abdominal ultrasound was normal, so a catheter was passed through the lumen and intestinal content was aspirated.

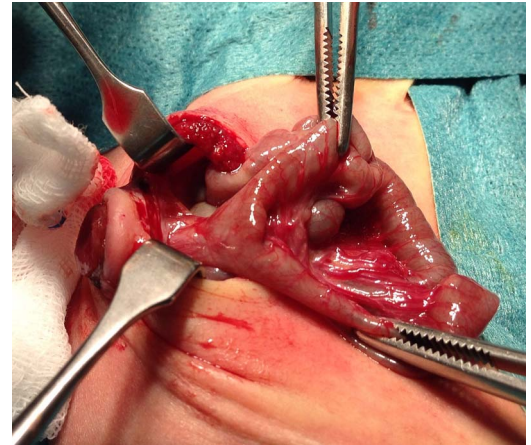
Surgical correction, with resection of the duct, took place on the fifth day of life (figure 2).

Histology revealed proximal intestinal mucosa consistent with patent omphalomesenteric duct (OMD). The newborn regained normal bowel function and resumed breastfeeding after 5 days.

The OMD is an embryological structure that connects the midgut to the yolk sac and normally regresses between the fifth and ninth weeks of fetal development.<sup>1</sup>

Remnants of the OMD may persist in approximately 2% of the population, the most common presentation being Meckel's diverticulum. Symptoms occur most frequently during childhood years and vary greatly, as patients may be asymptomatic, have abdominal pain, painless rectal bleeding, or intussusception/prolapse of ileum at the umbilicus, among others.<sup>1-3</sup>

An accurate diagnosis and management of this condition, with a highly unusual presentation, is important as it may lead to invagination of the



**Figure 2** Surgery demonstrating the patent omphalomesenteric duct.

small bowel and intestinal obstruction, with a significant increase in mortality.<sup>2</sup>

## Learning points

- ▶ A patent omphalomesenteric duct, although rare, may be easily identifiable in the newborn period, such as in an incidental finding after sectioning the umbilical cord.
- ▶ The differential diagnosis is mainly with patent urachus and umbilical polyp.
- ▶ If there is doubt about the nature of the umbilical discharge, an ultrasound or fistulogram can contribute to a final diagnosis and delineate the anatomy before surgery, however, if the discharge is obviously faecal, there is no need for investigative techniques.



**Figure 1** Umbilical cord with two umbilical arteries, one umbilical vein and a fourth structure with a lumen discharging intestinal content.

**Contributors** IRP was involved in data acquisition and drafting of the manuscript. TM and SL participated in the drafting and revising of the manuscript. MF participated in the conception and revising of the manuscript.

**Competing interests** None.

**Patient consent** Obtained.

**Provenance and peer review** Not commissioned; externally peer reviewed.

## REFERENCES

- 1 Tamilselvan K, Mohan A, Cheslyn-Curtis S, *et al.* Persistent umbilical discharge from an omphalomesenteric duct cyst containing gastric mucosa. *Case Rep Pediatr* 2012;2012:482185.
- 2 Pauleau G, Commandeur D, Andro C, *et al.* Intestinal prolapse through a persistente omphalomesenteric duct causing small-bowel obstruction. *S Afr J Surg* 2012;50:102-3.
- 3 Moore TC. Omphalomesenteric duct malformations. *Semin Pediatr Surg* 1996;5:116-23.



CrossMark

**To cite:** Periquito IR, Marques T, Lima S, *et al.* *BMJ Case Rep* Published online: [please include Day Month Year] doi:10.1136/bcr-2014-206553

Copyright 2014 BMJ Publishing Group. All rights reserved. For permission to reuse any of this content visit <http://group.bmj.com/group/rights-licensing/permissions>.  
BMJ Case Report Fellows may re-use this article for personal use and teaching without any further permission.

Become a Fellow of BMJ Case Reports today and you can:

- ▶ Submit as many cases as you like
- ▶ Enjoy fast sympathetic peer review and rapid publication of accepted articles
- ▶ Access all the published articles
- ▶ Re-use any of the published material for personal use and teaching without further permission

For information on Institutional Fellowships contact [consortiasales@bmjgroup.com](mailto:consortiasales@bmjgroup.com)

Visit [casereports.bmj.com](http://casereports.bmj.com) for more articles like this and to become a Fellow